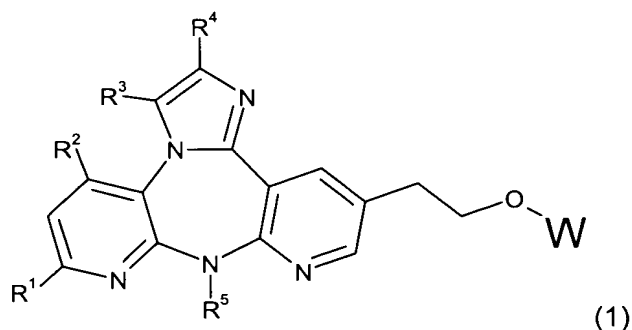


# ABSTRACT

Compounds represented by formula 1:

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wherein **R**<sup>1</sup> is H, halogen, (C<sub>1-4</sub>)alkyl, O(C<sub>1-4</sub>)alkyl, and haloalkyl; **R**<sup>2</sup> is H or methyl; **R**<sup>3</sup> is H or (C<sub>1-4</sub>)alkyl; **R**<sup>4</sup> is H or (C<sub>1-4</sub>)alkyl; **R**<sup>5</sup> is (C<sub>1-4</sub>)alkyl, (C<sub>1-4</sub>)alkyl(C<sub>3-7</sub>)cyclo-alkyl or (C<sub>3-7</sub>)cycloalkyl; and **W** is a fused phenyl-5 or 6-membered heterocycle having one or two heteroatoms selected from N or S; or **W** is phenyl, 1,1'-biphenyl, 2, 3-dihydro-1*H*-indene, 1, 2, 3, 4-tetrahydronaphthyl, or naphthyl; said **W** being optionally substituted with (C<sub>1-4</sub>)alkyl, which in turn can be optionally substituted with a carboxy or (C<sub>1-4</sub>)alkoxycarbonyl, or a salt or ester thereof. The compounds have inhibitory activity against Wild Type, single and double mutant strains of HIV.

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